

STN Columbus

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NEWS 2 NOV 21 CAS patent coverage to include exemplified prophetic
substances identified in English-, French-, German-,
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NEWS 15 FEB 11 WTEXTILES reloaded and enhanced
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NEWS 20 FEB 23 TOXCENTER updates mirror those of MEDLINE - more
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* * * * * STN Columbus * * * * *

FILE 'HOME' ENTERED AT 22:22:24 ON 13 MAR 2009

=> file uspatall

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	1.54	1.54

FILE 'USPATFULL' ENTERED AT 22:26:23 ON 13 MAR 2009
CA INDEXING COPYRIGHT (C) 2009 AMERICAN CHEMICAL SOCIETY (ACS)

FILE 'USPATOLD' ENTERED AT 22:26:23 ON 13 MAR 2009
CA INDEXING COPYRIGHT (C) 2009 AMERICAN CHEMICAL SOCIETY (ACS)

FILE 'USPAT2' ENTERED AT 22:26:23 ON 13 MAR 2009
CA INDEXING COPYRIGHT (C) 2009 AMERICAN CHEMICAL SOCIETY (ACS)

=> s tamsulosin
L1 1288 TAMSULOSIN

=> s tamsulosin/clm
L2 226 TAMSULOSIN/CLM

=> s polymer?
L3 1224605 POLYMER?

=> s polymer?/clm
L4 432182 POLYMER?/CLM

=> s (calcium or potassium or sodium or magnesium)
L5 1287207 (CALCIUM OR POTASSIUM OR SODIUM OR MAGNESIUM)

=> s (calcium or potassium or sodium or magnesium)/clm
L6 305082 (CALCIUM OR POTASSIUM OR SODIUM OR MAGNESIUM)/CLM

=> s (multilayer or layer)
L7 2063906 (MULTILAYER OR LAYER)

=> s (multilayer or layer)/clm
L8 810896 (MULTILAYER OR LAYER)/CLM

=> s capsule?
L9 243183 CAPSULE?

=> s capsule?/clm
L10 27197 CAPSULE?/CLM

=> s l1 and l4
L11 242 L1 AND L4

=> s l1 and l3
L12 897 L1 AND L3

=> s l5 and l12
L13 888 L5 AND L12

=> s l7 and l13
L14 600 L7 AND L13

=> s l9 and l14
L15 493 L9 AND L14

=> s l2 and l4
L16 50 L2 AND L4

=> s l6 and l16
L17 33 L6 AND L16

=> s l8 and l17
L18 6 L8 AND L17

=> s l10 and l18
L19 2 L10 AND L18

=> d 1-2

L19 ANSWER 1 OF 2 USPATFULL on STN

Full Text

AN 2009:4272 USPATFULL
TI Controlled release tamsulosin hydrochloride formulation
IN Cheng, Xiu Xiu, Weston, FL, UNITED STATES
Cheng, Xiufang, Weston, FL, UNITED STATES
PA Watson Pharmaceuticals, Inc. (U.S. corporation)
PI US 20090004284 A1 20090101
AI US 2007-821990 A1 20070626 (11)
DT Utility
FS APPLICATION
LN.CNT 567
INCL INCLM: 424/497.000
INCLS: 424/490.000; 514/603.000
NCL NCLM: 424/497.000
NCLS: 424/490.000; 514/603.000
IC IPCI A61K0009-14 [I,A]; A61K0031-18 [I,A]
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L19 ANSWER 2 OF 2 USPATFULL on STN

Full Text

AN 2008:253184 USPATFULL
TI Advanced drug development and manufacturing
IN Birnbaum, Eva R., Los Alamos, NM, UNITED STATES
Koppisch, Andrew T., Flagstaff, AZ, UNITED STATES
Baldwin, Sharon M., Santa Fe, NM, UNITED STATES
Warner, Benjamin P., Los Alamos, NM, UNITED STATES
McCleskey, T. Mark, Los Alamos, NM, UNITED STATES
Stewart, Jeffrey Joseph, Los Alamos, NM, UNITED STATES
Berger, Jennifer A., Los Alamos, NM, UNITED STATES
Harris, Michael N., Los Alamos, NM, UNITED STATES
Burrell, Anthony K., Los Alamos, NM, UNITED STATES
PI US 20080220441 A1 20080911
AI US 2007-974156 A1 20071010 (11)
RLI Continuation-in-part of Ser. No. US 2001-859701, filed on 16 May 2001,
PENDING Continuation-in-part of Ser. No. US 2002-206524, filed on 25 Jul
2002, ABANDONED Continuation-in-part of Ser. No. US 2003-621825, filed
on 16 Jul 2003, Pat. No. US 6858148
PRAI US 2006-850594P 20061010 (60)
DT Utility
FS APPLICATION
LN.CNT 10199
INCL INCLM: 435/071.000
INCLS: 436/501.000; 436/172.000; 436/086.000; 378/045.000
NCL NCLM: 435/007.100
NCLS: 378/045.000; 436/086.000; 436/172.000; 436/501.000
IC IPCI G01N0033-53 [I,A]; G01N0021-76 [I,A]; G01N0033-68 [I,A];
G01N0023-223 [I,A]; G01N0023-22 [I,C*]
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

=> d 118 1-6

L18 ANSWER 1 OF 6 USPATFULL on STN

Full Text

AN 2009:4272 USPATFULL
TI Controlled release tamsulosin hydrochloride formulation
IN Cheng, Xiu Xiu, Weston, FL, UNITED STATES
Cheng, Xiufang, Weston, FL, UNITED STATES
PA Watson Pharmaceuticals, Inc. (U.S. corporation)
PI US 20090004284 A1 20090101
AI US 2007-821990 A1 20070626 (11)
DT Utility
FS APPLICATION
LN.CNT 567
INCL INCLM: 424/497.000
INCLS: 424/490.000; 514/603.000
NCL NCLM: 424/497.000
NCLS: 424/490.000; 514/603.000
IC IPCI A61K0009-14 [I,A]; A61K0031-18 [I,A]
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L18 ANSWER 2 OF 6 USPATFULL on STN

Full Text

AN 2008:354338 USPATFULL
TI Solid form
IN Darmuzey, Olivia, Brussels, BELGIUM
MacLeod, Graeme, Wezembeek Oppem, BELGIUM
Cengic, Dzenana, Brussels, BELGIUM
PI US 20080311162 A1 20081218
AI US 2007-803825 A1 20070516 (11)
DT Utility
FS APPLICATION
LN.CNT 1512
INCL INCLM: 424/401.000
INCLS: 424/490.000; 514/263.340
NCL NCLM: 424/401.000
NCLS: 424/490.000; 514/263.340
IC IPCI A61K0008-02 [I,A]; A61K0009-14 [I,A]; A61K0031-522 [I,A];
A61K0031-519 [I,C*]; C11D0017-06 [I,A]
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L18 ANSWER 3 OF 6 USPATFULL on STN

Full Text

AN 2008:268489 USPATFULL
TI UROLOGICAL MEDICAL DEVICES FOR RELEASE OF THERAPEUTIC AGENTS
IN Cheng, Eric, Bloomington, IN, UNITED STATES
Li, Jianmin, Lexington, MA, UNITED STATES
Bucay-Couto, Weena, Burlington, MA, UNITED STATES
Sanders, Scott, Hinsdale, IL, UNITED STATES
Schuermann, James F., Natick, MA, UNITED STATES
Sheu, Min-Shyan, Chelmsford, MA, UNITED STATES
PA Boston Scientific Scimed, Inc., Maple Grove, MN, UNITED STATES (U.S.
corporation)
PI US 20080234659 A1 20080925
AI US 2008-52037 A1 20080320 (12)
PRAI US 2007-919081P 20070320 (60)
DT Utility
FS APPLICATION
LN.CNT 1180
INCL INCLM: 604/523.000
INCLS: 623/023.660
NCL NCLM: 604/523.000
NCLS: 623/023.660
IC IPCI A61M0025-00 [I,A]; A61F0002-04 [I,A]
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L18 ANSWER 4 OF 6 USPATFULL on STN

Full Text

AN 2008:253184 USPATFULL
TI Advanced drug development and manufacturing
IN Birnbaum, Eva R., Los Alamos, NM, UNITED STATES
Koppisch, Andrew T., Flagstaff, AZ, UNITED STATES
Baldwin, Sharon M., Santa Fe, NM, UNITED STATES
Warner, Benjamin P., Los Alamos, NM, UNITED STATES
McCleskey, T. Mark, Los Alamos, NM, UNITED STATES
Stewart, Jeffrey Joseph, Los Alamos, NM, UNITED STATES
Berger, Jennifer A., Los Alamos, NM, UNITED STATES
Harris, Michael N., Los Alamos, NM, UNITED STATES
Burrell, Anthony K., Los Alamos, NM, UNITED STATES
PI US 20080220441 A1 20080911
AI US 2007-974156 A1 20071010 (11)
RLI Continuation-in-part of Ser. No. US 2001-859701, filed on 16 May 2001,
PENDING Continuation-in-part of Ser. No. US 2002-206524, filed on 25 Jul
2002, ABANDONED Continuation-in-part of Ser. No. US 2003-621825, filed
on 16 Jul 2003, Pat. No. US 6858148
PRAI US 2006-850594P 20061010 (60)
DT Utility
FS APPLICATION
LN.CNT 10199
INCL INCLM: 435/071.000
INCLS: 436/501.000; 436/172.000; 436/086.000; 378/045.000
NCL NCLM: 435/007.100
NCLS: 378/045.000; 436/086.000; 436/172.000; 436/501.000

IC IPCI G01N0033-53 [I,A]; G01N0021-76 [I,A]; G01N0033-68 [I,A];
G01N0023-223 [I,A]; G01N0023-22 [I,C*]
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L18 ANSWER 5 OF 6 USPATFULL on STN

Full Text

AN 2008:86578 USPATFULL
TI Tamsulosin controlled-release tablet
IN Gan, Yong, Huairou, CHINA
Zhou, Xinteng, Huairou, CHINA
PA Ocean Star International, Inc., Snowville, UT, UNITED STATES (non-U.S.
corporation)
PI US 20080075775 A1 20080327
AI US 2006-580215 A1 20061011 (11)
PRAI CN 2006-10153091 20060922
DT Utility
FS APPLICATION
LN.CNT 720
INCL INCLM: 424/473.000
NCL NCLM: 424/473.000
IC IPCI A61K0009-24 [I,A]
IPCR A61K0009-24 [I,C]; A61K0009-24 [I,A]
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L18 ANSWER 6 OF 6 USPATFULL on STN

Full Text

AN 2007:140489 USPATFULL
TI Sustained release formulations
IN Cho, Seong Hwan, Suwon-city, KOREA, REPUBLIC OF
Ku, Jeong, Yongin-city, KOREA, REPUBLIC OF
Lim, Dong Kwon, Yongin-city, KOREA, REPUBLIC OF
Cheon, Jun Hee, Suwon-city, KOREA, REPUBLIC OF
An, Tae Kun, Yongin-city, KOREA, REPUBLIC OF
Ko, Jae Kyoung, Incheon-city, KOREA, REPUBLIC OF
Youn, Yong Sik, Yongin-city, KOREA, REPUBLIC OF
Park, Choong Sil, Icheon-city, KOREA, REPUBLIC OF
Suh, Hea Ran, Icheon-city, KOREA, REPUBLIC OF
Yang, Eun Young, Suwon-city, KOREA, REPUBLIC OF
Jeon, Eun Kyung, Yongin-city, KOREA, REPUBLIC OF
Kim, Chang Ju, Suwon-city, KOREA, REPUBLIC OF
PA CJ CORPORATION, Seoul, KOREA, REPUBLIC OF, 100-749 (non-U.S.
corporation)
PI US 20070122480 A1 20070531
AI US 2004-574337 A1 20040925 (10)
WO 2004-KR2496 20040925
20060509 PCT 371 date
PRAI KR 2003-67588 20030929
KR 2004-77158 20040924
DT Utility
FS APPLICATION
LN.CNT 799
INCL INCLM: 424/472.000
NCL NCLM: 424/472.000
IC IPCI A61K0009-24 [I,A]
IPCR A61K0009-24 [I,C]; A61K0009-24 [I,A]; A61K0009-22 [I,C*];
A61K0009-22 [I,A]
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

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(FILE 'HOME' ENTERED AT 22:22:24 ON 13 MAR 2009)

FILE 'USPATFULL, USPATOLD, USPAT2' ENTERED AT 22:26:23 ON 13 MAR 2009

L1 1288 S TAMSULOSIN
L2 226 S TAMSULOSIN/CLM
L3 1224605 S POLYMER?
L4 432182 S POLYMER?/CLM
L5 1287207 S (CALCIUM OR POTASSIUM OR SODIUM OR MAGNESIUM)
L6 305082 S (CALCIUM OR POTASSIUM OR SODIUM OR MAGNESIUM)/CLM
L7 2063906 S (MULTILAYER OR LAYER)
L8 810896 S (MULTILAYER OR LAYER)/CLM

L9 243183 S CAPSULE?
 L10 27197 S CAPSULE?/CLM
 L11 242 S L1 AND L4
 L12 897 S L1 AND L3
 L13 888 S L5 AND L12
 L14 600 S L7 AND L13
 L15 493 S L9 AND L14
 L16 50 S L2 AND L4
 L17 33 S L6 AND L16
 L18 6 S L8 AND L17
 L19 2 S L10 AND L18

=> d 115 483-493

L15 ANSWER 483 OF 493 USPAT2 on STN

Full Text

AN 2003:100119 USPAT2
 TI Quinazolone derivatives as alpha 1A/B adrenergic receptor antagonists
 IN Becker, Cyrus Kephra, Menlo Park, CA, UNITED STATES
 Caroon, Joan Marie, Mountain View, CA, UNITED STATES
 Melville, Chris Richard, Palo Alto, CA, UNITED STATES
 Padilla, Fernando, Fremont, CA, UNITED STATES
 Pfister, Jurg Roland, Los Altos, CA, UNITED STATES
 Zhang, Xiaoming, Campbell, CA, UNITED STATES
 PA Syntex (U.S.A.) LLC, Palo Alto, CA, UNITED STATES (U.S. corporation)
 PI US 6900220 B2 20050531
 AI US 2002-40319 20020102 (10)
 PRAI US 2001-259337P 20010102 (60)
 US 2001-325267P 20010927 (60)
 DT Utility
 FS GRANTED
 LN.CNT 2798
 INCL INCLM: 514/266.210
 INCLS: 514/234.200; 514/234.500; 514/249.000; 514/252.170; 514/264.100;
 514/266.200; 544/116.000; 544/117.000; 544/279.000; 544/284.000;
 544/350.000
 NCL NCLM: 514/266.210; 514/223.200
 NCLS: 514/234.200; 514/234.500; 514/249.000; 514/252.170; 514/264.100;
 514/266.200; 544/116.000; 544/117.000; 544/279.000; 544/284.000;
 544/350.000; 544/012.000
 IC [7]
 ICM A61K031-517
 ICS A61K031-535; C07D487-00; C07D417-00; C07D471-00
 IPCI C07D0285-22 [ICM,7]; C07D0285-00 [ICM,7,C*]; A61K0031-549
 [ICS,7]; A61K0031-517 [ICS,7]
 IPCI-2 A61K0031-517 [ICM,7]; A61K0031-535 [ICS,7]; C07D0487-00 [ICS,7];
 C07D0417-00 [ICS,7]; C07D0471-00 [ICS,7]
 IPCR A61K0031-517 [I,C*]; A61K0031-517 [I,A]; A61K0031-519 [I,C*];
 A61K0031-519 [I,A]; A61K0031-549 [I,C*]; A61K0031-549 [I,A];
 C07D0401-00 [I,C*]; C07D0401-04 [I,A]; C07D0401-14 [I,A];
 C07D0403-00 [I,C*]; C07D0403-14 [I,A]; C07D0405-00 [I,C*];
 C07D0405-14 [I,A]; C07D0417-00 [I,C*]; C07D0417-04 [I,A];
 C07D0417-14 [I,A]; C07D0471-00 [I,C*]; C07D0471-04 [I,A];
 C07D0471-14 [I,A]; C07D0487-00 [I,C*]; C07D0487-04 [I,A];
 C07D0491-00 [I,C*]; C07D0491-04 [I,A]
 EXF 514/234.2; 514/234.5; 514/249; 514/252.17; 514/264.1; 514/266.2;
 514/266.21; 544/116; 544/117; 544/279; 544/284; 544/350
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L15 ANSWER 484 OF 493 USPAT2 on STN

Full Text

AN 2003:92739 USPAT2
 TI Solid carriers for improved delivery of hydrophobic active ingredients
 in pharmaceutical compositions
 IN Patel, Mahesh V., Salt Lake City, UT, United States
 Chen, Feng-Jing, Salt Lake City, UT, United States
 PA Lipocine, Inc., Salt Lake City, UT, United States (U.S. corporation)
 PI US 6569463 B2 20030527
 AI US 2001-800593 20010306 (9)
 RLI Division of Ser. No. US 1999-447690, filed on 23 Nov 1999, now patented,
 Pat. No. US 6248363
 DT Utility

FS GRANTED
 LN.CNT 3198
 INCL INCLM: 424/497.000
 INCLS: 424/422.000; 424/427.000; 424/430.000; 424/433.000; 424/434.000;
 424/435.000; 424/436.000; 424/441.000; 424/451.000; 424/457.000;
 424/463.000; 424/464.000; 424/465.000; 424/466.000; 424/470.000;
 424/474.000; 424/476.000; 424/482.000; 424/490.000; 424/489.000;
 424/498.000; 514/773.000; 514/784.000; 514/785.000; 514/786.000;
 514/779.000
 NCL NCLM: 424/497.000; 424/465.000
 NCLS: 424/422.000; 424/427.000; 424/430.000; 424/433.000; 424/434.000;
 424/435.000; 424/436.000; 424/441.000; 424/451.000; 424/457.000;
 424/463.000; 424/464.000; 424/465.000; 424/466.000; 424/470.000;
 424/474.000; 424/476.000; 424/482.000; 424/489.000; 424/490.000;
 424/498.000; 514/773.000; 514/779.000; 514/784.000; 514/785.000;
 514/786.000; 977/906.000; 977/927.000
 IC [7]
 ICM A61K009-16
 ICS A61K009-28; A61K009-32; A61K009-52; A61K009-56; A61K009-58
 IPCI A61K0009-20 [ICM, 7]; A61K0009-16 [ICS, 7]; A61K0009-50 [ICS, 7]
 IPCI-2 A61K0009-16 [ICM, 7]; A61K0009-28 [ICS, 7]; A61K0009-32 [ICS, 7];
 A61K0009-30 [ICS, 7, C*]; A61K0009-52 [ICS, 7]; A61K0009-56 [ICS, 7];
 A61K0009-58 [ICS, 7]; A61K0009-52 [ICS, 7, C*]
 IPCR A61K0009-02 [I, C*]; A61K0009-02 [I, A]; A61K0009-14 [I, C*];
 A61K0009-14 [I, A]; A61K0009-16 [I, C*]; A61K0009-16 [I, A];
 A61K0009-20 [I, C*]; A61K0009-20 [I, A]; A61K0009-30 [I, C*];
 A61K0009-32 [I, A]; A61K0009-48 [I, C*]; A61K0009-48 [I, A];
 A61K0009-50 [I, C*]; A61K0009-50 [I, A]; A61K0009-51 [I, C*];
 A61K0009-51 [I, A]; A61K0009-52 [I, C*]; A61K0009-52 [I, A];
 A61K0009-54 [I, A]; A61K0009-56 [I, A]; A61K0009-58 [I, A];
 A61K0031-21 [I, C*]; A61K0031-216 [I, A]; A61K0031-232 [I, A];
 A61K0031-351 [I, C*]; A61K0031-351 [I, A]; A61K0031-366 [I, C*];
 A61K0031-366 [I, A]; A61K0031-40 [I, C*]; A61K0031-40 [I, A];
 A61K0031-403 [I, C*]; A61K0031-404 [I, A]; A61K0031-415 [I, C*];
 A61K0031-415 [I, A]; A61K0031-4196 [I, C*]; A61K0031-4196 [I, A];
 A61K0031-421 [I, C*]; A61K0031-421 [I, A]; A61K0031-4353 [I, C*];
 A61K0031-436 [I, A]; A61K0031-4409 [I, C*]; A61K0031-4409 [I, A];
 A61K0031-4427 [I, C*]; A61K0031-4439 [I, A]; A61K0031-472 [I, C*];
 A61K0031-4725 [I, A]; A61K0031-519 [I, C*]; A61K0031-522 [I, A];
 A61K0031-57 [I, C*]; A61K0031-57 [I, A]; A61K0031-64 [I, C*];
 A61K0031-64 [I, A]; A61K0031-662 [I, C*]; A61K0031-663 [I, A];
 A61K0038-23 [I, C*]; A61K0038-23 [I, A]; A61K0047-02 [I, C*];
 A61K0047-02 [I, A]; A61K0047-10 [I, C*]; A61K0047-10 [I, A];
 A61K0047-14 [I, C*]; A61K0047-14 [I, A]; A61K0047-22 [I, C*];
 A61K0047-22 [I, A]; A61K0047-26 [I, C*]; A61K0047-26 [I, A];
 A61K0047-28 [I, C*]; A61K0047-28 [I, A]; A61K0047-32 [I, C*];
 A61K0047-32 [I, A]; A61K0047-36 [I, C*]; A61K0047-36 [I, A];
 A61K0047-38 [I, C*]; A61K0047-38 [I, A]; A61K0047-44 [I, C*];
 A61K0047-44 [I, A]; A61P0001-00 [I, C*]; A61P0001-04 [I, A];
 A61P0003-00 [I, C*]; A61P0003-04 [I, A]; A61P0003-06 [I, A];
 A61P0003-10 [I, A]; A61P0005-00 [I, C*]; A61P0005-16 [I, A];
 A61P0005-24 [I, A]; A61P0005-40 [I, A]; A61P0007-00 [I, C*];
 A61P0007-02 [I, A]; A61P0007-10 [I, A]; A61P0009-00 [I, C*];
 A61P0009-04 [I, A]; A61P0009-06 [I, A]; A61P0009-10 [I, A];
 A61P0009-12 [I, A]; A61P0013-00 [I, C*]; A61P0013-08 [I, A];
 A61P0015-00 [I, C*]; A61P0015-10 [I, A]; A61P0017-00 [I, C*];
 A61P0017-12 [I, A]; A61P0019-00 [I, C*]; A61P0019-06 [I, A];
 A61P0019-10 [I, A]; A61P0021-00 [I, C*]; A61P0021-02 [I, A];
 A61P0025-00 [I, C*]; A61P0025-04 [I, A]; A61P0025-06 [I, A];
 A61P0025-08 [I, A]; A61P0025-16 [I, A]; A61P0025-20 [I, A];
 A61P0025-22 [I, A]; A61P0025-26 [I, A]; A61P0025-28 [I, A];
 A61P0029-00 [I, C*]; A61P0029-00 [I, A]; A61P0031-00 [I, C*];
 A61P0031-04 [I, A]; A61P0031-10 [I, A]; A61P0031-12 [I, A];
 A61P0033-00 [I, C*]; A61P0033-06 [I, A]; A61P0033-10 [I, A];
 A61P0035-00 [I, C*]; A61P0035-00 [I, A]; A61P0037-00 [I, C*];
 A61P0037-06 [I, A]; A61P0043-00 [I, C*]; A61P0043-00 [I, A]
 EXF 424/422; 424/433; 424/436; 424/435; 424/440; 424/451; 424/452; 424/464;
 424/465; 424/482; 424/489; 424/490; 424/480; 424/463; 424/470; 424/497;
 424/498; 424/476; 424/427; 424/430; 424/434; 424/441; 424/466; 424/474
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.

Full Text

AN 2003:78121 USPAT2
TI Modified release formulations containing a hypnotic agent
IN Lemmens, Jacobus M., Mook, NETHERLANDS
van den Heuvel, Dennie J. M., Boxmeer, NETHERLANDS
Platteeuw, Johannes J., s'Hertogenbosch, NETHERLANDS
van Dalen, Frans, Nijmegen, NETHERLANDS
PA Synthon BV, Nijmegen, NETHERLANDS (non-U.S. corporation)
PI US 6638535 B2 20031028
AI US 2001-833662 20010413 (9)
PRAI US 2000-196939P 20000413 (60)
DT Utility
FS GRANTED
LN.CNT 902
INCL INCLM: 424/489.000
INCLS: 424/490.000; 424/464.000; 424/465.000; 514/300.000; 514/781.000
NCL NCLM: 424/489.000
NCLS: 424/464.000; 424/465.000; 424/490.000; 514/300.000; 514/781.000
IC [7]
ICM A61K009-20
ICS A61K009-14; A61K009-16; A61K031-44; A61K047-00
IPCI A61K0009-14 [ICM,7]
IPCI-2 A61K0009-20 [ICM,7]; A61K0009-14 [ICS,7]; A61K0009-16 [ICS,7];
A61K0031-44 [ICS,7]; A61K0047-00 [ICS,7]
IPCR A61K0009-16 [I,C*]; A61K0009-16 [I,A]; A61K0009-26 [I,C*];
A61K0009-26 [I,A]; A61K0031-4353 [I,C*]; A61K0031-437 [I,A];
A61K0031-519 [I,C*]; A61K0031-519 [I,A]
EXF 424/489; 424/490; 424/464; 424/465; 514/300; 514/781
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L15 ANSWER 486 OF 493 USPAT2 on STN

Full Text

AN 2002:308400 USPAT2
TI Heterocyclic compounds their production and use
IN Tasaka, Akihiro, Suita, JAPAN
Hitaka, Takenori, Takarazuka, JAPAN
Matsutani, Etsuya, Suita, JAPAN
PA Takeda Chemical Industries, Ltd., Osaka, JAPAN (non-U.S. corporation)
PI US 6716863 B2 20040406
WO 2001077107 20011018
AI US 2001-889974 20010724 (9)
WO 2001-JP2937 20010405
PRAI JP 2000-106836 20000407
DT Utility
FS GRANTED
LN.CNT 3144
INCL INCLM: 514/374.000
INCLS: 548/235.000
NCL NCLM: 514/374.000
NCLS: 548/235.000
IC [7]
ICM A61K031-422
ICS C07D413-12
IPCI C07D0413-02 [ICM,7]; C07D0413-00 [ICM,7,C*]; A61K0031-422 [ICS,7]
IPCI-2 A61K0031-422 [ICM,7]; C07D0413-12 [ICS,7]; C07D0413-00 [ICS,7,C*]
IPCR A61P0035-00 [I,C*]; A61P0035-00 [I,A]; C07D0263-00 [I,C*];
C07D0263-32 [I,A]; C07D0413-00 [I,C*]; C07D0413-12 [I,A]
EXF 514/374; 548/235
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L15 ANSWER 487 OF 493 USPAT2 on STN

Full Text

AN 2002:301622 USPAT2
TI Benzimidazoles that are useful in treating male sexual dysfunction
IN Cowart, Marlon D., Round Lake Beach, IL, UNITED STATES
Bhatia, Pramila A., Libertyville, IL, UNITED STATES
Daanen, Jerome F., Racine, WI, UNITED STATES
Stewart, Andrew O., Libertyville, IL, UNITED STATES
Patel, Meena V., Green Oaks, IL, UNITED STATES
Kolasa, Teodozyj, Lake Villa, IL, UNITED STATES
Brioni, Jorge D., Vernon Hills, IL, UNITED STATES
Rohde, Jeffrey, Evanston, IL, UNITED STATES

PA Abbott Laboratories, Abbott Park, IL, UNITED STATES (U.S. corporation)
 PI US 7022728 B2 20060404
 AI US 2002-94265 20020308 (10)
 PRAI US 2001-340452P 20011214 (60)
 US 2001-296078P 20010605 (60)
 US 2001-274805P 20010309 (60)
 DT Utility
 FS GRANTED
 LN.CNT 2990
 INCL INCLM: 514/395.000
 INCLS: 514/394.000; 514/393.000; 514/252.140; 514/254.060; 514/253.010
 NCL NCLM: 514/395.000; 514/252.190
 NCLS: 514/252.140; 514/253.010; 514/254.060; 514/393.000; 514/394.000;
 514/253.090; 514/254.030
 IC IPCI A61K0031-496 [ICM, 7]
 IPCI-2 A61K0031-415 [I,A]; A61K0031-495 [I,A]; A61K0031-50 [I,A]
 IPCR A61K0031-496 [I,C*]; A61K0031-496 [I,A]; A61K0031-415 [I,A];
 A61K0031-415 [I,C]; A61K0031-495 [I,C]; A61K0031-495 [I,A];
 A61K0031-50 [I,C]; A61K0031-50 [I,A]
 EXF 514/255; 514/258; 514/394; 514/393; 514/359; 514/362; 514/363; 514/385;
 514/395; 514/252.14; 514/254.06; 514/253.01
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L15 ANSWER 488 OF 493 USPAT2 on STN

Full Text

AN 2002:288146 USPAT2
 TI As-needed administration of tricyclic and other non-SRI antidepressant
 drugs to treat premature ejaculation
 IN Tam, Peter, Redwood City, CA, UNITED STATES
 Gesundheit, Neil, Los Altos, CA, UNITED STATES
 Wilson, Leland F., Menlo Park, CA, UNITED STATES
 PA Vivus, Inc., Mountain View, CA, UNITED STATES (U.S. corporation)
 PI US 6946141 B2 20050920
 AI US 2001-996407 20011121 (9)
 RLI Continuation-in-part of Ser. No. US 2000-721412, filed on 21 Nov 2000,
 Pat. No. US 6495154
 DT Utility
 FS GRANTED
 LN.CNT 1495
 INCL INCLM: 424/423.000
 INCLS: 424/434.000; 424/435.000; 424/443.000; 424/449.000; 424/451.000;
 424/464.000; 424/045.000; 424/046.000
 NCL NCLM: 424/423.000; 514/278.000
 NCLS: 424/045.000; 424/046.000; 424/434.000; 424/435.000; 424/443.000;
 424/449.000; 424/451.000; 424/464.000
 IC [7]
 ICM A61F002-02
 ICS A61F013-02; A61K009-48; A61K009-70; A61K009-04
 IPCI A61K0031-44 [ICM, 7]
 IPCI-2 A61F0002-02 [ICM, 7]; A61F0013-02 [ICS, 7]; A61K0009-48 [ICS, 7];
 A61K0009-70 [ICS, 7]; A61K0009-04 [ICS, 7]
 IPCR A61K0009-02 [I,C*]; A61K0009-02 [I,A]; A61K0009-08 [I,C*];
 A61K0009-08 [I,A]; A61K0009-12 [I,C*]; A61K0009-12 [I,A];
 A61K0009-19 [I,C*]; A61K0009-19 [I,A]; A61K0009-20 [I,C*];
 A61K0009-20 [I,A]; A61K0009-46 [I,C*]; A61K0009-46 [I,A];
 A61K0031-135 [I,C*]; A61K0031-135 [I,A]; A61K0031-136 [I,C*];
 A61K0031-136 [I,A]; A61K0031-403 [I,C*]; A61K0031-404 [I,A];
 A61K0031-55 [I,C*]; A61K0031-55 [I,A]; A61K0031-551 [I,C*];
 A61K0031-551 [I,A]; A61K0031-553 [I,C*]; A61K0031-553 [I,A];
 A61K0031-554 [I,C*]; A61K0031-554 [I,A]; A61K0045-00 [I,C*];
 A61K0045-00 [I,A]; A61K0045-06 [I,A]; A61K0047-34 [I,C*];
 A61K0047-34 [I,A]; A61P0015-00 [I,C*]; A61P0015-00 [I,A];
 A61P0043-00 [I,C*]; A61P0043-00 [I,A]
 EXF 424/423; 424/434; 424/435; 424/443; 424/449; 424/451; 424/464; 424/45;
 424/46
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L15 ANSWER 489 OF 493 USPAT2 on STN

Full Text

AN 2002:67175 USPAT2
 TI Administration of phosphodiesterase inhibitors for the treatment of
 premature ejaculation

IN Wilson, Leland F., Menlo Park, CA, United States
Doherty, Jr., Paul C., Cupertino, CA, United States
Place, Virgil A., Kawaihae, HI, United States
Smith, William L., Montclair, NJ, United States
Abdel-Hamid Abdou Ali, Ibrahim AbouBakr, Mansoura, EGYPT
PA Vivus, Inc., Mountain View, CA, United States (U.S. corporation)
PI US 6403597 B2 20020611
AI US 2001-888250 20010621 (9)
RLI Continuation-in-part of Ser. No. US 1999-467094, filed on 10 Dec 1999
Continuation-in-part of Ser. No. US 1998-181070, filed on 27 Oct 1998,
now patented, Pat. No. US 6037346, issued on 14 Mar 2000
Continuation-in-part of Ser. No. US 1997-958816, filed on 28 Oct 1997,
now abandoned
DT Utility
FS GRANTED
LN.CNT 2030
INCL INCLM: 514/256.000
NCL NCLM: 514/256.000; 514/001.000
IC [7]
ICM A61K031-50
IPCI A61K0031-00 [ICM,7]
IPCI-2 A61K0031-50 [ICM,7]
IPCR A61K0009-00 [I,C*]; A61K0009-00 [I,A]; A61K0031-00 [I,C*];
A61K0031-00 [I,A]; A61K0031-343 [I,C*]; A61K0031-343 [I,A];
A61K0031-381 [I,C*]; A61K0031-381 [I,A]; A61K0031-40 [I,C*];
A61K0031-40 [I,A]; A61K0031-4015 [I,C*]; A61K0031-4015 [I,A];
A61K0031-4164 [I,C*]; A61K0031-4164 [I,A]; A61K0031-4166 [I,A];
A61K0031-426 [I,C*]; A61K0031-426 [I,A]; A61K0031-4353 [I,C*];
A61K0031-437 [I,A]; A61K0031-4427 [I,C*]; A61K0031-4439 [I,A];
A61K0031-444 [I,A]; A61K0031-4704 [I,C*]; A61K0031-4704 [I,A];
A61K0031-4709 [I,C*]; A61K0031-4709 [I,A]; A61K0031-4738 [I,C*];
A61K0031-4745 [I,A]; A61K0031-496 [I,C*]; A61K0031-496 [I,A];
A61K0031-50 [I,C*]; A61K0031-50 [I,A]; A61K0031-502 [I,C*];
A61K0031-502 [I,A]; A61K0031-5025 [I,C*]; A61K0031-5025 [I,A];
A61K0031-505 [I,C*]; A61K0031-505 [I,A]; A61K0031-519 [I,C*];
A61K0031-519 [I,A]; A61K0031-52 [I,A]; A61K0031-522 [I,A];
A61K0031-5375 [I,C*]; A61K0031-538 [I,A]; A61K0031-549 [I,C*];
A61K0031-549 [I,A]; A61K0031-551 [I,C*]; A61K0031-5513 [I,A];
A61K0045-00 [I,C*]; A61K0045-06 [I,A]
EXF 514/258
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L15 ANSWER 490 OF 493 USPAT2 on STN

Full Text

AN 2002:55008 USPAT2
TI Clear oil-containing pharmaceutical compositions containing a
therapeutic agent
IN Chen, Feng-Jing, Salt Lake City, UT, United States
Patel, Mahesh V., Salt Lake City, UT, United States
Fikstad, David T., Salt Lake City, UT, United States
PA Lipocine, Inc., Salt Lake City, UT, United States (U.S. corporation)
PI US 6761903 B2 20040713
AI US 2001-877541 20010608 (9)
RLI Continuation-in-part of Ser. No. US 1999-345615, filed on 30 Jun 1999,
now patented, Pat. No. US 6267985 Continuation-in-part of Ser. No. US
2000-751968, filed on 29 Dec 2000, now patented, Pat. No. US 6458383
Continuation-in-part of Ser. No. US 1999-375636, filed on 17 Aug 1999,
now patented, Pat. No. US 6309663
DT Utility
FS GRANTED
LN.CNT 3614
INCL INCLM: 424/451.000
INCLS: 424/043.000; 424/433.000; 424/436.000; 424/441.000; 424/445.000;
424/455.000; 424/456.000; 424/458.000; 424/463.000; 424/464.000;
424/465.000; 424/489.000; 424/490.000; 424/725.000; 514/772.200;
514/772.300; 514/777.000; 514/779.000; 514/781.000; 514/783.000;
514/784.000; 514/785.000; 514/786.000; 514/937.000; 514/944.000
NCL NCLM: 424/451.000; 514/054.000
NCLS: 424/043.000; 424/433.000; 424/436.000; 424/441.000; 424/445.000;
424/455.000; 424/456.000; 424/458.000; 424/463.000; 424/464.000;
424/465.000; 424/489.000; 424/490.000; 424/725.000; 514/772.200;
514/772.300; 514/777.000; 514/779.000; 514/781.000; 514/783.000;

514/784.000; 514/785.000; 514/786.000; 514/937.000; 514/944.000;
 424/727.000; 424/731.000; 424/750.000; 424/757.000

IC [7]
 ICM A61K009-08
 ICS A61K009-10; A61K009-14; A61K009-20; A61K009-48
 IPCI A61K0031-715 [ICM,7]; A61K0035-78 [ICS,7]
 IPCI-2 A61K0009-08 [ICM,7]; A61K0009-10 [ICS,7]; A61K0009-14 [ICS,7];
 A61K0009-20 [ICS,7]; A61K0009-48 [ICS,7]
 IPCR A61K0009-48 [I,C*]; A61K0009-48 [I,A]; A61K0036-185 [I,C*];
 A61K0036-47 [I,A]

EXF 424/451; 424/450; 424/433; 424/436; 424/441; 424/443; 424/445
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L15 ANSWER 491 OF 493 USPAT2 on STN

Full Text

AN 2002:27516 USPAT2
 TI Compounds and methods to increase plasma HDL cholesterol levels and
 improve HDL functionality
 IN Luchoomun, Jayraz, Lilburn, GA, UNITED STATES
 Meng, Charles Q., Alpharetta, GA, UNITED STATES
 Saxena, Uday, Atlanta, GA, UNITED STATES
 Sikorski, James A., Alpharetta, GA, UNITED STATES
 PA Atherogenics, Inc., Alpharetta, GA, UNITED STATES (U.S. corporation)
 PI US 6881860 B2 20050419
 AI US 2001-833407 20010411 (9)
 PRAI US 2000-196201P 20000411 (60)
 DT Utility
 FS GRANTED
 LN.CNT 3107
 INCL INCLM: 562/426.000
 INCLS: 514/568.000
 NCL NCLM: 562/426.000; 514/517.000
 NCLS: 514/571.000; 514/649.000; 558/037.000; 564/347.000

IC [7]
 ICM C07C321-00
 IPCI A61K0031-255 [ICM,7]; A61K0031-21 [ICM,7,C*]; A61K0031-192
 [ICS,7]; A61K0031-185 [ICS,7,C*]; A61K0031-145 [ICS,7]
 IPCI-2 C07C0321-00 [ICM,7]
 IPCR C07C0323-00 [I,C*]; C07C0323-20 [I,A]; G01N0033-92 [I,C*];
 G01N0033-92 [I,A]

EXF 562/426; 514/568
 CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L15 ANSWER 492 OF 493 USPAT2 on STN

Full Text

AN 2002:21845 USPAT2
 TI Compositions and methods for improved delivery of hydrophobic agents
 IN Patel, Mahesh V., Salt Lake City, UT, United States
 Chen, Feng-Jing, Salt Lake City, UT, United States
 PA Lipocine, Inc., Salt Lake City, UT, United States (U.S. corporation)
 PI US 6451339 B2 20020917
 AI US 2001-898553 20010702 (9)
 RLI Continuation of Ser. No. US 1999-258654, filed on 26 Feb 1999, now
 patented, Pat. No. US 6294192
 DT Utility
 FS GRANTED
 LN.CNT 2907
 INCL INCLM: 424/451.000
 INCLS: 424/450.000; 424/455.000; 424/456.000; 424/463.000; 424/489.000;
 424/499.000; 424/502.000; 424/435.000; 424/464.000; 424/937.000;
 424/938.000; 424/939.000; 514/940.000; 514/941.000; 514/942.000;
 514/943.000; 514/975.000
 NCL NCLM: 424/451.000; 424/400.000
 NCLS: 424/435.000; 424/450.000; 424/455.000; 424/456.000; 424/463.000;
 424/464.000; 424/489.000; 424/499.000; 424/502.000; 514/937.000;
 514/938.000; 514/939.000; 514/940.000; 514/941.000; 514/942.000;
 514/943.000; 514/975.000

IC [7]
 ICM A61K009-127
 IPCI A61K0009-00 [ICM,7]
 IPCI-2 A61K0009-127 [ICM,7]
 IPCR A61K0009-48 [I,C*]; A61K0009-48 [I,A]; A61K0031-57 [I,C*];

A61K0031-57 [I,A]; A61K0038-12 [I,C*]; A61K0038-13 [I,A]
EXF 424/450; 424/451; 424/455; 424/456; 424/463; 424/489; 424/499; 424/502;
424/435; 424/464; 514/937; 514/938; 514/939; 514/940; 514/941; 514/943;
514/975

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L15 ANSWER 493 OF 493 USPAT2 on STN

Full Text

AN 2001:229703 USPAT2
TI Co-administration of a prostaglandin and an androgenic agent in the
treatment of female sexual dysfunction
IN Place, Virgil A., Kawaihae, HI, United States
Wilson, Leland F., Menlo Park, CA, United States
Doherty, Jr., Paul C., Cupertino, CA, United States
Hanamoto, Mark S., Belmont, CA, United States
Spivack, Alfred P., Menlo Park, CA, United States
Gesundheit, Neil, Los Altos, CA, United States
Bennett, Sean R., Denver, CO, United States
PA Vivus, Inc., Mountain View, CA, United States (U.S. corporation)
PI US 6593313 B2 20030715
AI US 2001-905458 20010713 (9)
RLI Continuation of Ser. No. US 2000-539484, filed on 30 Mar 2000, now
patented, Pat. No. US 6306841 Continuation of Ser. No. US 1998-181316,
filed on 27 Oct 1998, now abandoned Continuation-in-part of Ser. No. US
1997-959064, filed on 28 Oct 1997, now patented, Pat. No. US 5877216
Continuation-in-part of Ser. No. US 1997-959057, filed on 28 Oct 1997,
now abandoned
DT Utility
FS GRANTED
LN.CNT 1331
INCL INCLM: 514/108.000
NCL NCLM: 514/108.000; 514/530.000
NCLS: 514/288.000; 514/573.000
IC [7]
ICM A61K031-19
ICS A61K031-557
IPCI A61K0031-5575 [ICM,7]; A61K0031-557 [ICM,7,C*]; A61K0031-48
[ICS,7]
IPCI-2 A61K0031-19 [ICM,7]; A61K0031-185 [ICM,7,C*]; A61K0031-557
[ICS,7]
IPCR A61K0009-00 [I,C*]; A61K0009-00 [I,A]; A61K0009-02 [N,C*];
A61K0009-02 [N,A]; A61K0031-00 [I,C*]; A61K0031-00 [I,A];
A61K0031-15 [I,C*]; A61K0031-15 [I,A]; A61K0031-21 [I,C*];
A61K0031-21 [I,A]; A61K0031-28 [I,C*]; A61K0031-295 [I,A];
A61K0031-48 [I,C*]; A61K0031-48 [I,A]; A61K0031-5375 [I,C*];
A61K0031-5377 [I,A]; A61K0031-557 [I,C*]; A61K0031-557 [I,A];
A61K0031-5575 [I,A]; A61K0031-5585 [I,A]; A61K0031-56 [I,C*];
A61K0031-56 [I,A]; A61K0045-00 [I,C*]; A61K0045-06 [I,A]

EXF 514/573

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

=> d 115 kwic 492

L15 ANSWER 492 OF 493 USPAT2 on STN

DETD The POE-POP block copolymers are a unique class of **polymeric**
surfactants. The unique structure of the surfactants, with hydrophilic
POE and hydrophobic POP moieties in well-defined ratios and positions,
provides. . . Synperonic PE series (ICI); Pluronic® series
(BASF), Emkalyx, Lutrol (BASF), Supronic, Monolan, Pluracare, and
Plurodac. The generic term for these **polymers** is "poloxamer" (CAS
9003-11-6). These **polymers** have the formula:
DETD . . . use in the present invention. Preferred anionic surfactants
include fatty acid salts and bile salts. Specifically, preferred ionic
surfactants include **sodium** oleate, **sodium** lauryl sulfate, **sodium**
lauryl sarcosinate, **sodium** dioctyl sulfosuccinate, **sodium** cholate,
and **sodium** taurocholate. Examples of such surfactants are shown in
Table 18 below. For simplicity, typical counterions are shown in the
entries. . . in the art, however, that any bioacceptable counterion
may be used. For example, although the fatty acids are shown as **sodium**
salts, other cation counterions can also be used, such as alkali metal
cations or ammonium. Unlike typical non-ionic surfactants, these. . .

DETD
TABLE 18

Ionic Surfactants
COMPOUND HLB

FATTY ACID SALTS >10

Sodium caproate
Sodium caprylate
Sodium caprate
Sodium laurate
Sodium myristate
Sodium myristolate
Sodium palmitate
Sodium palmitoleate
Sodium oleate 18
Sodium ricinoleate
Sodium linoleate
Sodium linolenate
Sodium stearate
Sodium lauryl sulfate (dodecyl) 40
Sodium tetradecyl sulfate
Sodium lauryl sarcosinate
Sodium dioctyl sulfosuccinate [**sodium** docusate (Cytec)]

BILE SALTS >10

Sodium cholate
Sodium taurocholate
Sodium glycocholate
Sodium deoxycholate
Sodium taurodeoxycholate
Sodium glycodeoxycholate
Sodium ursodeoxycholate
Sodium chenodeoxycholate
Sodium taurochenodeoxycholate
Sodium glyco cheno deoxycholate
Sodium cholylsarcosinate
Sodium N-methyl taurocholate

PHOSPHOLIPIDS

Egg/Soy lecithin [Epikuron .TM. (Lucas Meyer), Ovothin .TM. (Lucas Meyer)]

Lyso egg/soy lecithin
Hydroxylated lecithin
Lysophosphatidylcholine
Cardiolipin
Sphingomyelin
Phosphatidylcholine
Phosphatidyl ethanolamine
Phosphatidic acid
Phosphatidyl glycerol
Phosphatidyl serine

PHOSPHORIC ACID ESTERS

Diethanolammonium polyoxyethylene-10. . . fatty alcohol ethoxylates
with phosphoric acid or anhydride

CARBOXYLATES

Ether carboxylates (by oxidation of terminal OH group of fatty
alcohol ethoxylates)
Succinylated monoglycerides [LAMEGIN ZE (Henkel)]
Sodium stearyl fumarate
Stearoyl propylene glycol hydrogen succinate
Mono/diacetylated tartaric acid esters of mono- and diglycerides
Citric acid esters of mono-, diglycerides
Glyceryl-lacto esters of fatty acids (CFR ref. 172.852)

Acyl lactylates:

lactylic esters of fatty acids
calcium/sodium stearyl-2-lactylate
calcium/sodium stearyl lactylate

Alginate salts

Propylene glycol alginate

SULFATES AND SULFONATES

Ethoxylated alkyl sulfates

Alkyl benzene sulfones

α -olefin sulfonates

Acyl isethionates
Acyl taurates
Alkyl glyceryl ether sulfonates
Octyl sulfosuccinate disodium
Disodium undecylenamideo-MEA-sulfosuccinate
CATIONIC. . .

DETD A third method of determining optical clarity and carrier diffusivity through the aqueous boundary **layer** is to quantitatively measure the size of the particles of which the dispersion is composed. These measurements can be performed. . .

DETD analgesics and anti-inflammatory agents, such as aloxiprin, auranofin, azapropazone, benorylate, capsaicin, celecoxib, diclofenac, diflunisal, etodolac, fenbufen, fenoprofen **calcium**, flurbiprofen, ibuprofen, indomethacin, ketoprofen, ketorolac, leflunomide, meclofenamic acid, mefenamic acid, nabumetone, naproxen, oxaprozin, oxyphenbutazone, phenylbutazone, piroxicam, refocoxib, sulindac, tetrahydrocannabinol, tramadol. . .

DETD anti-epileptics, such as beclamide, carbamazepine, clonazepam, ethotoin, felbamate, fosphenytoin **sodium**, lamotrigine, methoin, methsuximide, methylphenobarbitone, oxcarbazepine, paramethadione, phenacemide, phenobarbitone, phenytoin, phensuximide, primidone, sulthiame, tiagabine HCl, topiramate, valproic acid, and vigabatrin;

DETD gastrointestinal agents, such as bisacodyl, cimetidine, cisapride, diphenoxylate HCl, domperidone, famotidine, lansoprazole, loperamide, mesalazine, nizatidine, omeprazole, ondansetron HCl, rabeprazole **sodium**, ranitidine HCl and sulphasalazine;

DETD muscle relaxants, such as dantrolene **sodium** and tizanidine HCl;

DETD and others, such as becaplermin, donepezil HCl, L-thyroxine, methoxsalen, verteporfrin, physostigmine, pyridostigmine, raloxifene HCl, sibutramine HCl, sildenafil citrate, tacrine, **tamsulosin** HCl, and tolterodine.

DETD . . . a semi-solid dispersion or a solid dispersion. If desired, the compositions may be encapsulated in a hard or soft gelatin **capsule**, a starch **capsule** or an enteric coated **capsule**. The term "enteric coated **capsule**" as used herein means a **capsule** coated with a coating resistant to acid; i.e., an acid resistant enteric coating. Although solubilizers are typically used to enhance. . . of a hydrophobic therapeutic agent, they may also render the compositions more suitable for encapsulation in hard or soft gelatin **capsules**. Thus, the use of a solubilizer such as those described above is particularly preferred in **capsule** dosage forms of the pharmaceutical compositions. If present, these solubilizers should be added in amounts sufficient to impart to the. . .

DETD . . . is mono-modal and narrow. This reduced and more uniform size enables more efficient drug transport through the intestinal aqueous boundary **layer**, and through the absorptive brush border membrane. More efficient transport to absorptive sites leads to improved and more consistent absorption. . .

DETD . . . conditions which limit production of lipase, such as pancreatic lipase secretory diseases; and dependence of lipolysis on stomach pH, endogenous **calcium** concentration, and presence of co-lipase or other digestion enzymes. The lack of lipolysis dependence further provides transport which does not. . .

DETD . . . the pharmaceutical compositions of the present invention allow for faster transport of the hydrophobic therapeutic agent through the aqueous boundary **layer**.

DETD . . . the hydrophobic therapeutic agent. A third solution was prepared with simulated intestinal fluid, plus an additional aliquot of 20 mM **sodium** taurocholate (a bile salt); this solution is designated SIFB in Table 29. Finally, a fourth solution was prepared with simulated intestinal fluid, 20 mM **sodium** taurocholate, and 5 mM lecithin; this solution is designated SIFBL. The 20 mM bile salt and 5 mM lecithin concentrations. . .

DETD

Cyclosporine 0.140 g
Cremophor RH-40 0.41 g
Arlacel 186 0.29 g
Sodium taurocholate 0.26 g
Propylene glycol 0.46 g

CLM What is claimed is:

. . . lecithins; lysolecithin and hydrogenated lysolecithins;

lysophospholipids and derivatives thereof; phospholipids and derivatives thereof; salts of alkylsulfates; salts of fatty acids; **sodium** docusate; and mixtures thereof.

CLM What is claimed is:

. . . and salts, analogues, and derivatives thereof; lecithins, lysolecithin, phospholipids, lysophospholipids and derivatives thereof; salts of alkylsulfates; salts of fatty acids; **sodium** docusate; acyl lactylates; monoacetylated tartaric acid esters of monoglycerides, monoacetylated tartaric acid esters of diglycerides, diacetylated tartaric acid esters of. . .

CLM What is claimed is:

77. A dosage form comprising a **capsule** filled with the pharmaceutical formulation of any one of claims 69, 70, 71 or 72.

CLM What is claimed is:

78. The dosage form of claim 77, wherein the **capsule** is a hard gelatin **capsule**, a soft gelatin **capsule**, a starch **capsule** or an enteric coated **capsule**.

CLM What is claimed is:

88. The formulation of claim 1, wherein the formulation is contained in a **capsule**.

CLM What is claimed is:

95. The formulation of claim 89, wherein the formulation is contained in a **capsule**.

CLM What is claimed is:

105. The formulation of claim 104, wherein the formulation is contained in a **capsule**.

CLM What is claimed is:

107. The formulation of claim 106, wherein the formulation is contained in a **capsule**.

CLM What is claimed is:

117. The formulation of claim 116, wherein the formulation is contained in a **capsule**.

=> log y

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

ENTRY

SESSION

FULL ESTIMATED COST

42.32

43.86

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